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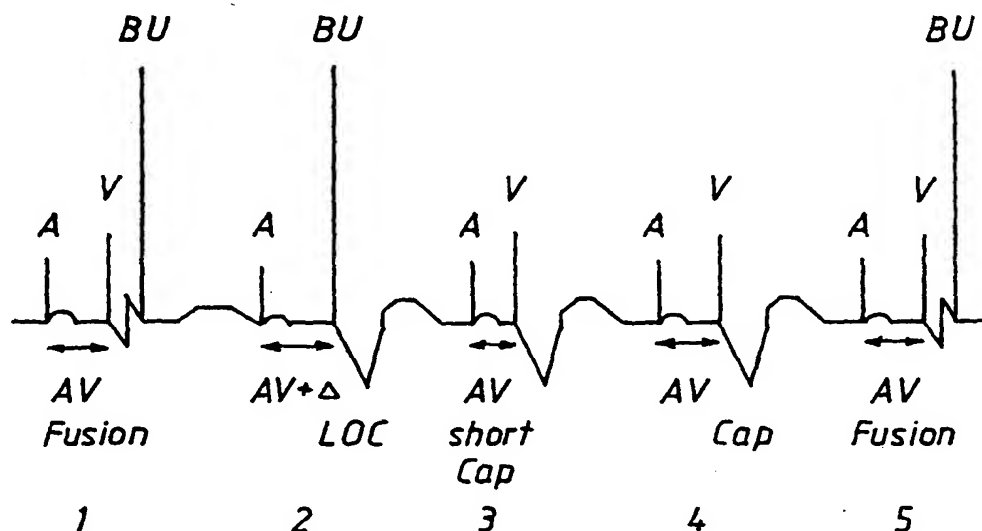
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(54) Title: IMPLANTABLE HEART STIMULATOR



(57) Abstract: Implantable heart stimulator comprising an AV-interval generator adapted to generate a variable AV-interval and provided with a predetermined basic AV-interval. The heart stimulator further comprises a counter that counts the number of times the AV-interval is changed in response of a fusion avoidance algorithm or a stimulation threshold search algorithm during a predetermined time period, e.g. 1-3 minutes, wherein the basic AV-interval is changed if said number of times is greater than a predetermined value, e.g. 4-10.

WO 01/03769 A1

Implantable heart stimulatorTechnical field of the invention.

The present invention relates to an implantable heart
5 stimulator according to the preamble of the independent
claim.

Background of the invention.

To reduce the energy consumption of heart stimulators, an
automatic threshold search function, is used to maintain the
10 energy of the stimulation pulses at a level just above that
which is needed to effectuate capture, cf. e.g. US-A-
5,458,623. A reliable detection of the evoked response,
which then is necessary, is, however, not a simple matter,
especially when it is desired to sense the evoked response
15 with the same electrode as the one delivering the
stimulation pulse and in particular if the sensing is
performed by a unipolar electrode configuration.

A fusion beat refers typically in pacing to the ECG waveform
which results when an intrinsic depolarisation and a
20 pacemaker output pulse occur simultaneously and both
contribute to the electrical activation of that heart
chamber. Closely related to a fusion beat is a pseudofusion
beat that refers to a spontaneous cardiac depolarisation
occurring at or near a pulse generator output pulse. Because
25 the stimulus occurs after the heart has spontaneously
depolarised, the pacemaker output is ineffective, but it
distorts the morphology of the complex on the ECG.

Today, fusion beats create a problem for the automatic
threshold search function since these beats often are not
30 detected as heart beats. Instead the heart stimulator in

question interprets the evoked response as a loss of capture and as a consequence a backup pulse is issued and the stimulation pulse amplitude is increased. Following undetected fusion beats the heart stimulator might deliver
5 back-up pulses (high output mode), until the next threshold search is performed. This misinterpretation by the heart stimulator of the evoked response signal will, of course, increase the current drain and decrease the lifetime of the battery and the automatic threshold search function will be
10 disabled for some time.

US-5,713,930 discloses a dual chamber pacing system and method with control of the AV interval. The AV interval is adjusted to provide for an optimal AV setting for a selected pacing application. A ventricular fusion test is performed,
15 wherein variations in QT interval are monitored corresponding to variations in AV interval. Based upon the AV-QT data, the pacemaker can determine the ventricular fusion zone where the pacemaker AV interval is substantially the same as the intrinsic conduction interval, as well as
20 the knee where AV intervals just shorter than the ventricular fusion zone result in full capture. One application of the dual chamber pacemaker disclosed in US-5,713,930 is for patients with intermittent AV conduction or occasional AV block where it is desired to set AV delay to
25 be just greater than the natural conduction interval, so that spontaneous beats are permitted.

It should be understood that the present invention is directed to an implantable heart stimulator having, inter alia, an inhibiting function, which means that if intrinsic
30 heart activity is detected, in the atrium or in the ventricle, no stimulation pulse is generated. This means that, using the commonly accepted terminology, the AV-interval could be started by an intrinsic atrial heart

activity, a P-wave, and the started interval is then a PV-interval. Thus, instead of writing PV/AV-interval is the term AV-interval used throughout this application.

With reference to figure 1 the behaviour of a fusion avoidance algorithm in a commercially available dual chamber heart stimulator now will be described. A dual chamber heart stimulator comprising a fusion avoidance algorithm and also a threshold search algorithm (discussed below) is disclosed in e.g. "User's manual for AFFINITY™DR, Model 5330 L/R, Dual-Chamber Pulse Generator with AUTOCAPTURE™ Pacing System", by Pacesetter, Ordering No. 2039782, Part. No 9192000-001, issued in 1998, pages 52-54.

Figure 1 shows an internal electrogram (IEGM) with a normal paced heartbeat seen as the first complex. "A" and "V" designate the stimulation pulses in the atrium and in the ventricle, respectively, and "Cap" stands for capture, i.e. the applied stimulation pulse in the ventricle was successful. "AV" designates the AV-interval. The next stimulation pulse applied in the ventricle did not result in capture and to ensure safe pacing a back-up pulse is applied a predetermined interval after the stimulation pulse. The loss of capture could be the result of an intrinsic contraction at the same time as the stimulation pulse is applied which is detected as a loss of capture. Another alternative is that the stimulation threshold for the heart tissue has increased.

In order to avoid fusion is the AV-interval prolonged with a predetermined time (in the figure designated as Δ), in other words, the heart stimulator prolongs the AV-interval and waits for an intrinsic activity. In this case, as can be seen in figure 1, a fusion beat was detected as loss of capture in spite it was an intrinsic beat. The AV-interval

is prolonged a predetermined number of times, e.g. 3-6 times.

If instead a loss of capture is the result of an increasing stimulation threshold an IEGM illustrating that case is shown in figure 2. The loss of capture (LOC) in the second complex is followed by a prolonged AV-interval $AV+\Delta$ in order to avoid a fusion beat as described above. In this case is the AV-interval $AV+\Delta$ timed out and a stimulation pulse (V) is applied to the ventricle. The applied stimulation pulse does not result in capture and a back-up pulse is applied which in turn results in capture. In this case was the loss of capture due to an increasing stimulation threshold of the heart tissue.

A threshold search algorithm may be activated by two consecutive loss of capture.

A preferred threshold search algorithm is illustrated in figure 2 where the AV-interval shortened to "AV-short" to override any intrinsic heart activity and the ventricular stimulation amplitude is successively stepped up by a predetermined amplitude step of e.g. 0,1-0,3 V and each unsuccessful ventricular stimulation pulse is followed by a back-up pulse. This is performed until the stimulation threshold is detected, i.e. capture is detected from the ventricular stimulation pulse, and the stimulation pulse amplitude is then set to a value that equals the stimulation threshold plus a working margin, e.g. 0,3 V.

The sinus node in the upper part of the atrium is the heart's own "pacemaker". In a normally functioning heart is a depolarisation wave generated by the sinus node and conducted along the conduction system from the atrium down to the ventricle. The conduction system is briefly heart

muscle cells specially adapted to depolarise at a certain frequency in the order of 0,9 - 2 Hz, corresponds to a heart rate of 54 - 120 beats per minute. If the conduction system from the sinus node in the atrium down to the ventricle does not work normally it is said that the patient has some kind of AV-block.

There are three major groups of AV-block.

A first degree AV-block is present if all atrial depolarisation are conducted to the ventricle but the PQ-interval is slightly prolonged (longer than 0,21s).

There are two types of second degree AV-blocks. In second degree AV-block, Mobitz type I (also called Wenchebach block), the PR-interval increases progressively until an impulse is not conducted to the ventricle. Thereafter the cycle is repeated again.

In second degree AV-block, Mobitz type II, the length of the PQ interval is usually stable, while the blocking pattern may be regular or irregular. This means that, if the blocking pattern is regular, every second up to seventh P-wave is blocked.

A third degree AV-block (complete block) is present when there is no conduction between the atria and ventricles.

Figure 3 shows an IEGM illustrating the behaviour of a dual chamber heart stimulator provided with a fusion avoidance algorithm in combination with a stimulation threshold search algorithm on a patient with a second degree AV-block, Mobitz type II.

The first complex represents a fusion beat followed by a back-up pulse. The AV-interval is then prolonged with Δ (see the fusion avoidance algorithm described above) in order to determine if the first complex was a fusion beat. In this case no intrinsic activity is detected in the ventricle

because the patient has a second degree AV-block, Mobitz type II (in the following called Mobitz II-block) and thus the prolonged AV-interval is timed out and a back-up pulse is generated. Since the criteria for initiating the stimulation threshold search algorithm is two consecutive loss of capture the search algorithm is thus initiated. The AV-interval is shortened to AV-short in order to override any intrinsic activity (see complex 3). The applied stimulation pulse was successful and capture was detected.

5 The threshold search algorithm is thereby terminated and the AV-interval is then restored to its original programmed or basic value (see complex 4). In the next complex (complex 5) another fusion beat is present and which is detected as a loss of capture. The AV-interval is prolonged again (as in

10 complex 2) and at the same time another Mobitz II-block is present and the sequence described above is repeated again.

15

One drawback with the above-described phenomena that occurs when fusion beats are present and in particular in patients with a Mobitz II-block is that some patients might feel unpleasant when the AV-interval constantly is changing.

20

Another drawback is that several high-energy back-up pulses are delivered which consumes energy.

Short description of the inventive concept.

25 The object of the present invention is to overcome the above-mentioned drawbacks and is achieved by an implantable heart stimulator according to the characterising portion of the independent claim.

Preferred embodiments are set forth in the dependant claims.

Short description of the appended drawings.

Figure 1 shows an IEGM illustrating a fusion avoidance algorithm used in the prior art;

Figure 2 shows an IEGM illustrating a stimulation threshold
5 search algorithm used in the prior art;

Figure 3 shows an IEGM illustrating the behaviour of a heart stimulator provided with a fusion avoidance algorithm in combination with a stimulation threshold search algorithm on a patient with a second degree AV-block, Mobitz type II;

10 Figure 4 shows a schematic block diagram of an implantable heart stimulator according to the invention.

Detailed description of preferred embodiments of the invention.

15 Figure 4 shows an implantable heart stimulator 2 comprising an atrial pulse generator 4, an atrial detector 6, a ventricular pulse generator 8 and a ventricular detector 10. The pulse generators 4,8 and the detectors 6,10 are adapted to be connected to one or many electrode leads arranged to
20 stimulate the heart tissue and/or to detect electrical heart activity in the atrium and in the ventricle, respectively.

A person skilled in the art is aware of many different ways to arrange the electrode lead or leads. According to one embodiment is two bipolar electrode leads arranged, whereas
25 one is arranged in the atrium and another in the ventricle. According to an alternative embodiment is one multipolar electrode lead with electrode surfaces arranged in the atrium for atrial stimulation/detection and electrode surfaces arranged in the ventricle for ventricular
30 stimulation/detection.

To be able to implement the invention in an implantable heart stimulator it must at least have the ability to sense electrical heart activity both in the atrium and in the ventricle and to stimulate in the ventricle.

- 5 The implantable heart stimulator further comprises a control unit 12 connected to the atrial and ventricular detectors 6,10, an AV interval generator 14 connected to the control unit 12 and a counter 16 connected to the AV interval generator. The AV interval generator 14 is also connected to
10 the atrial generator 4 and to the ventricular generator 8.

Upon detection of a P-wave or when the atrial stimulator generates an atrial stimulation pulse the AV interval generator starts an AV interval. The ventricular stimulator generates a ventricular stimulation pulse if the AV interval
15 is timed out. The ventricular stimulation can be inhibited if an intrinsic ventricular event is detected prior the expiration of the AV interval.

The control unit 12 includes means, e.g. a microprocessor, to activate different algorithms related to the operation of
20 the heart stimulator. Among these algorithms are the above mentioned stimulation threshold search algorithm and fusion avoidance algorithm.

The atrial and ventricular detectors are adapted to detect heart events in the atrium and in the ventricle,
25 respectively. The detection may be performed in many different ways. A specific steepness of the slope of the detected electrical heart signal could be used as an indicator of a specific heart event or a predetermined integration value of a predetermined interval following an
30 applied stimulation could also be used. The skilled person is aware of many other ways to perform the detection.

When a heart event is detected by any of the detectors a signal is applied to the control unit that judges *inter alia* whether the detected heart event was an intrinsic heart event or if it was a stimulated heart event.

- 5 The counter 16 connected to the AV interval generator 14 counts the number of times the AV interval (AVI) is changed during a predetermined time period. This can be performed in many different ways and one way is that each time the AV interval generator makes the ventricular stimulation
- 10 generator generate a ventricular stimulation pulse it compares the present AV interval (the AVI that just timed out) with the previous AV interval. If these consecutive AV intervals not are the same a count signal is generated and applied to the counter that counts the number of count
- 15 signals during a predetermined time period, e.g. during the last 1-3 minutes, preferably during the last 2 minutes. If this number is greater than a predetermined value the counter 16 generates an output signal which is applied to the AV interval generator causing the basic AV interval to
- 20 be changed. The predetermined value is in the interval 2-10, preferably 6. The basic AV interval is prolonged or shortened by a time in the interval of ± 30 ms, preferably ± 20 ms, if the number is greater than the predetermined value. In some cases it can be interesting to count even
- 25 fewer AVI changes during a shorter time period, e.g. if the predetermined value is 1 then if 2 changes occur during the last one minute (or parts of a minute) the basic AVI should be changed.

- A prerequisite for counting the number of times that the AVI
- 30 is changed is that the AVI is timed out and as a consequence of that a ventricular stimulation pulse is generated.

A person skilled in the art of heart stimulation is aware of many different situations where the AVI can be changed. In e.g. rate responsive heart stimulators the AVI can be controlled by a signal representing the detected activity.

- 5 The higher activity the shorter AVI is a relationship that is applied. It must be observed that the changes of the AVI of interest in the present application are only those that are related to the fusion avoidance algorithm and to the stimulation threshold search algorithm.
- 10 According to a preferred embodiment of the invention is the basic AVI prolonged with 20 ms if the number of changes is greater than a predetermined value during the last 2 minutes. If the situation (predetermined number of changes of the new basic AVI) is repeated again during a following
- 15 time period, 1-10 minutes, the basic AVI is shortened by 20 ms. Other sequences of the changes are of course possible, e.g. first prolong the AVI with 20 ms, then go back to the original basic AVI and then shorten the AVI by 20 ms.

- The present invention is not limited to the above-described
- 20 preferred embodiments. Various alternatives, modifications and equivalents may be used. Therefore, the above embodiments should not be taken as limiting the scope of the invention, which is defined by the appendant claims.

Claims

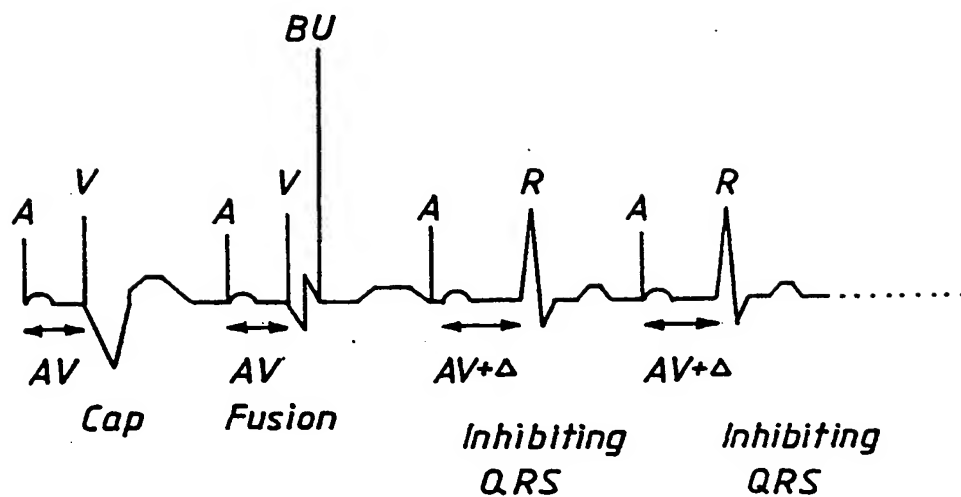
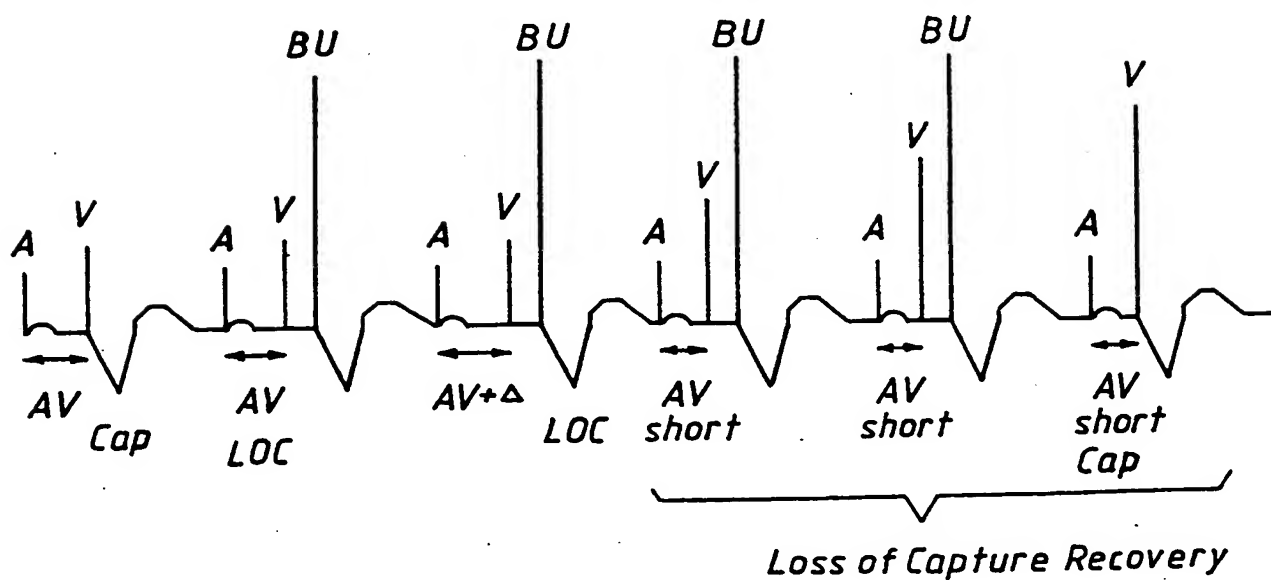
1. Implantable heart stimulator (2) comprising an AV-
interval generator (14) adapted to generate a variable AV-
5 interval and provided with a predetermined basic AV-
interval, characterized in that the heart stimulator
(2) further comprises a counter (16) that counts the number
of times the AV-interval is changed during a predetermined
time period, and where said counter (16) generates and
10 applies an output signal to said AV-signal generator (14) to
change said basic AV-interval if said number of times is
greater than a predetermined value.
2. Heart stimulator according to claim 1
15 characterized in that the change of the basic AV-
interval is in the interval -30 to 30 milliseconds.
3. Heart stimulator according to claim 2
characterized in that the change of the basic AV-
20 interval is -20 ms or +20 ms.
4. Heart stimulator according to any preceding claim
characterized in that said predetermined value is in
the range 2-10.
25
5. Heart stimulator according to any preceding claim
characterized in that said predetermined time period
is between 1 and 3 minutes.
- 30 6. Heart stimulator according to any preceding claim
characterized in that said heart stimulator further
comprises a control unit (12) provided with a fusion
avoidance algorithm.

7. Heart stimulator according to claim 6
characterized in that said fusion avoidance
algorithm, when activated, makes the AV-interval generator
5 (14) temporarily prolong the AV-interval.

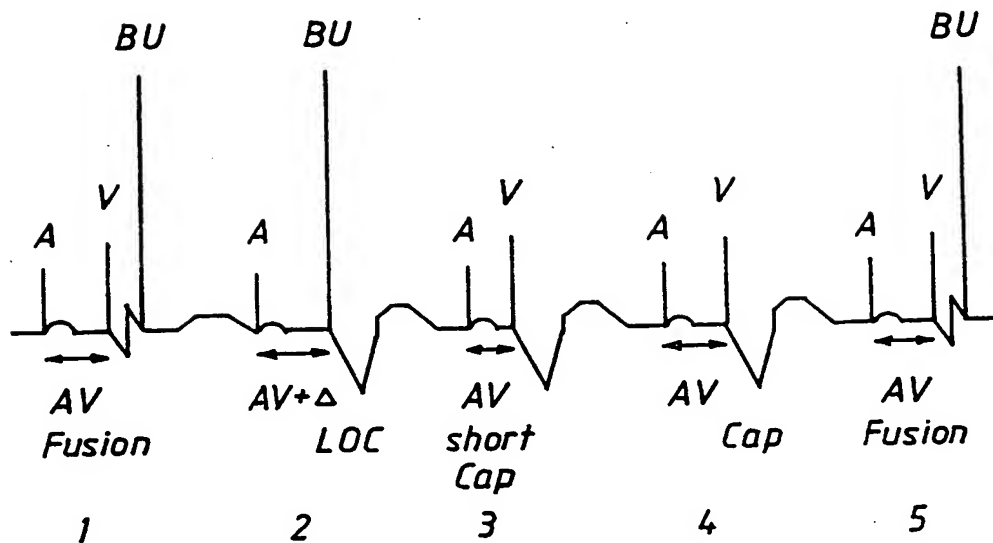
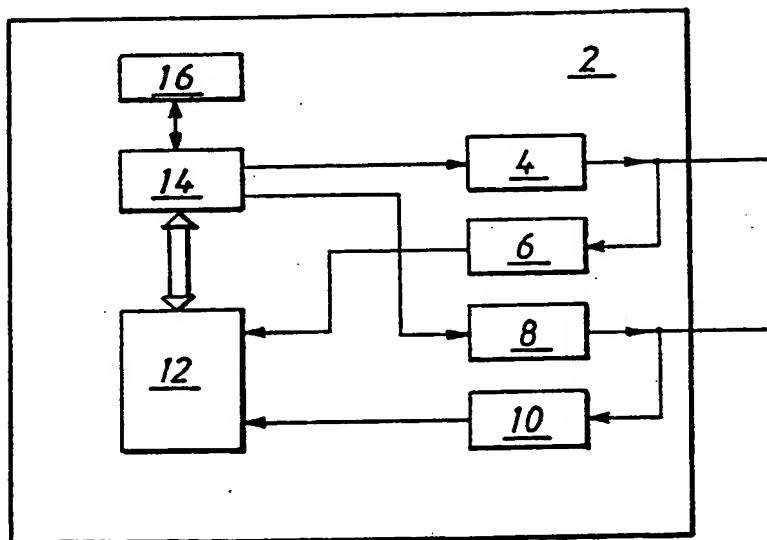
8. Heart stimulator according to any preceding claim
characterized in that said heart stimulator further
comprises a control unit (12) provided with a stimulation
10 threshold search algorithm.

9. Heart stimulator according to claim 8
characterized in that said stimulation threshold
search algorithm, when activated, makes the AV-interval
15 generator (14) temporarily shorten the AV-interval.

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Fig. 1Fig. 2

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Fig. 3Fig. 4

INTERNATIONAL SEARCH REPORT

International application No.

PCT/SE 00/01306

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: A61N 1/362

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: A61N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPODOC

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 4211235 A (JOHN W. KELLER, JR. ET AL), 8 July 1980 (08.07.80) --	1-9
A	US 5330511 A (WILLEM BOUTE), 19 July 1994 (19.07.94), abstract --	1-9
A	US 5340361 A (JASON A. SHOLDER), 23 August 1994 (23.08.94), abstract --	1-9
A	US 5683428 A (PER FRANBERG), 4 November 1997 (04.11.97), abstract -- -----	1-9

☐ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

* Special categories of cited documents:

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"&" document member of the same patent family

Date of the actual completion of the international search

25 Sept 2000

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

01/08/00

PCT/SE 00/01306

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